1. Anti-neoplastic and/or anti-leukemic effective compound selected from the formulae A and B:

wherein

for general formula A: wherein R_1 is mono or dihalogenated acyl group, aroyl group (Table 1), alkyloxy-carbonyl group or aryloxy-carbonyl group (Table 2) and R_3 is hydrogen or halogenated group, and R_2 is hydrogen or acetyl groups; wherein R_4 is PhCO or Me COCO or $CH_3CH=C(CH_3CH=(CH_3)CO, R_3$ is a halogenated group (Tables 1 and 2);

for general formula B: wherein R_1 is mono or dihalogenated acyl group or aroyl group (Table 1), alkyloxy-carbonyl group or aryloxy-carbonyl group (Table 2) and R_2 is hydrogen or acetyl group, and R_5 is any group from Table 3; R_6 is H or Me;

and wherein groups of Tables 1, 2 and 3 are selected from

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Table 1 Structures of Halogenated Acyl and Aroyl Groups

O _I
, Appr
0 مماليمة
x
(
, br.
£
) - pr

X: halogen (Cl'or Br or I or F)

Group 22 X	Group 29	X X	Group 35	X O pr
Group 23	Group 30	X	Group 36	X ₁
Group 25	Group 31	X X	Group 37	X ₂
Group 26 X	Group 32	X N Jord	Group 38	X O O O
Grou 27 X	Group 33	X V V	Group 39	×
Group 28 X O	Group 34	X O Y		x. o
✓ x			Group 40	N Jr

X: halogen (Cl or Br or I or F)
X₁: one type of halogen
X₂: other type of halogen

Table 2 Structures of Halogenated Alkyloxy- and Aryloxy- Carbonyl Groups

				arbonyl Group	os	
Group 41	×~~°~	Group 48	CX of	Group 55	1 0	
1	X OH OH	}		1	Y '0'	_z z
Group 43	НО X О	Group 50	X CLOY,	Group 57	ÓMe ,×	
					x-{Sloly	
1	X N L		ł		1	
	X JOHAT					
Group 47	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	X ₂ Group 54	X ₂ X ₁	Group 61	Sign	
X: halos	gen (Clor Bror Lo	T				

X: halogen (Cl or Br or I or F)
X₁: one type of halogen
X₂: other type of halogen

Table 2 (Contd)

Group 62	$X_{2} \longrightarrow R \bigcirc X_{1}$	Group 68	X ₂ 0	Group 74	x S S J
Group 63	X_2 X_1 X_1	Group69	X OH O J	Group 75	× S o
Group 64	X X O Y	Group70	x~~o ¹ / ₂ / ₂ / ₃	Group 76	x s pr
Group 65	× × × × ×	Group71	~ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	Group77	N O O O O O
Group 66	× \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	Group72	N N N O N O N O N O N O N O N O N O N O	Group78	× O N Pr
X₂ Group67	X10 X1	Group73	X N N N N N N N N N N N N N N N N N N N	Group79	× N O J'

X: halogen (Cl or Br or I or F)
X₁: one type of halogen
X₂: other type of halogen

Table 2 (Contd)

Group 80	× John	Group 86		Group 91	
	Х	l	X OLY	1	i
Group 82	× ° ° × ° ° × ° × ° × ° × ° × ° × ° × °	Group 88	× ° , , , , , , , , ,	Group 93	× Coly
Group 83	x o i , t	Group 89	R O O	Group 94	× o d
Group 84	x Doly	Group 90	X oly	Group 95	X N O I
Group 85	X O O				X S S S S

X: halogen (Cl or Br or I or F)

Table 3. Group Structures of Amino Acids and Their Codes Used in This Patent

Sec.

2. A compound of claim 1 of the formula:

TYPE I

wherein R_1 is a group selected from Table 1 (groups 1 to 40) and R_2 is H or $Ac_{\mathbf{p}}$

3. A compound of claim 1 of the formula:

TYPE II

wherein R_1 is a group selected from Table 2 (groups 41 to 95); R_2 is H or Ac

4. A compound of claim 1 of the formula:

TYPE III

wherein R_3 is a group selected from Table 1 (groups 1 to 40); and R_2 is H or Ac, and R_4 is PhCO or Me₃COCO or CH₃CH=C(CH₃)CO $_{\odot}$

5. A compound of claim 1 of the formula:

TYPE IV

wherein R_3 is a group selected from Table 2, (groups 41 to 95), R_2 is Ac or H, and R_4 is PhCO or Me₃COCO or CH₃CH=C(CH₃)CO₄

6. A compound of claim 1 of the formula:

TYPE V

R₂ is H or Ac;

R₃ is a group selected from Table 2 (groups 41 to 95)

7. A compound of claim 1 of the formula:

TYPE VI

wherein R_1 is a group selected from Table 2 (groups 41 to 95);

R₂ is H or Ac;

R₃ is a group selected from Table 1 (groups 1 to 40).

TYPE VII

 \widetilde{OR}_3 \widetilde{OH}_2 \widetilde{H} \widetilde{OAc} wherein R_1 is a group selected from Table 1 (groups 1 to 40); R_2 is H or Ac;

R₃ is a group selected from Table 1 (groups 1 to 40);

9. A compound of claim 1 of the formula:

TYPE VIII

OH H OAC OBZ wherein R₁ is a group from Table 2 (groups 41 to 95);

R₂ is H or Ac;

R₃ is a group selected from Table 2 (groups 41 to 95)

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A compound of claim 1 of the formula: 10.

TYPE IX

wherein R₁ is a group selected from Table 1 (groups 1 to 40);

R2 is H or Ac;

 R_5 is H or Me or G_1 or G_2 or G_3 or G_4 or G_5 or G_6 or G_7 or G_8 or G_9 or G_{11} or G_{12} or G₁₃;

R₆ is H, only in the case when R₅ is G₁₀ the group R₆ is H or Me₄

11. A compound of claim 1 of the formula:

TYPE X

wherein R₁ is a group selected from Table 2 (groups 55 to 95);

R₂ is H or Ac;

 R_5 is H or Me or G_1 or G_2 or G_3 or G_4 or G_5 or G_6 or G_7 or G_8 or G_9 or G_{11} or G_{12} or G_{13} ;

 R_6 is H, only in the case when R_5 is G_{10} the group R_6 is H or Meà

- 12. A pharmaceutical formulation which comprises as an active ingredient a compound of claim 1 or a pharmaceutically acceptable salt thereof.
- 13. A pharmaceutical formulation which comprises as an active ingredient a compound of claim 2 or a pharmaceutically acceptable salt thereof.
- 14. A pharmaceutical formulation which comprises as an active ingredient a compound of claim 3 or a pharmaceutically acceptable salt thereof.
- 15. A pharmaceutical formulation which comprises as an active ingredient a compound of claim 4 or a pharmaceutically acceptable salt thereof.
- 16. A pharmaceutical formulation which comprises as an active ingredient a compound of claim 5 or a pharmaceutically acceptable salt thereof.
- 17. A pharmaceutical formulation which comprises as an active ingredient a compound of claim 6 or a pharmaceutically acceptable salt thereof.
- 18. A pharmaceutical formulation which comprises as an active ingredient a compound of claim 7 or a pharmaceutically acceptable salt thereof.
- 19. A pharmaceutical formulation which comprises as an active ingredient a compound of claim 8 or a pharmaceutically acceptable salt thereof.
- 20. A pharmaceutical formulation which comprises as an active ingredient a compound of claim 9 or a pharmaceutically acceptable salt thereof.
- 21. A pharmaceutical formulation which comprises as an active ingredient a compound of claim 10 or a pharmaceutically acceptable salt thereof.

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- A pharmaceutical formulation which comprises as an active ingredient a 22. compound of claim 11 or a pharmaceutically acceptable salt thereof.
- A method for treating humans in need thereof comprising administering 23. to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 1.
- A method for treating humans in need thereof comprising administering 24. to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 2.
- A method for treating humans in need thereof comprising administering 25. to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 3.
- A method for treating humans in need thereof comprising administering 26. to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 4.
- A method for treating humans in need thereof comprising administering 27. to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 5.
- A method for treating humans in need thereof comprising administering 28. to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 6.
- 29. A method for treating humans in need thereof comprising administering to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 7.
- A method for treating humans in need thereof comprising administering 30. to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 8.
- A method for treating humans in need thereof comprising administering 31. to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 9.

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- 32. A method for treating humans in need thereof comprising administering to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 10.
- 33. A method for treating humans in need thereof comprising administering to said humans an anti-cancer or anti-leukemic effective amount of the compound of claim 11.